In the claims:

1. (Currently Amended) A compound of formula (I)

$$R^5$$
 R^4
 R^3
 R^2
 R^7
 R^1
 R^1

X is CH₂ or SO₂

R¹ is an optionally substituted aryl;

R² is carboxy, cyano, -C(O)CH₂OH, -CONHR⁸, -SO₂NHR⁹, tetrazol 5-yl, or SO₃H where R⁸ is selected from hydrogen, alkyl, aryl, cyano, hydroxy, -SO₂R¹² where R¹² is alkyl, aryl, heteroaryl, or haloalkyl, or R⁸ is a group (CHR¹³), -COOH where r is an integer of 1-3 and each R¹³ group is independently selected from hydrogen or alkyl; R⁹ is hydrogen, alkyl, optionally substituted aryl such as optionally substituted phenyl or optionally substituted_heteroaryl such as 5 or 6 membered heteroaryl groups, or a group COR¹⁴ where R¹⁴ is alkyl, aryl, heteroaryl or haloalkyl;

- R³ is hydrogen, a functional group, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted alkoxy, optionally substituted aralkyl, optionally substituted aralkyloxy or_optionally substituted eyeloalkyl;
- R^4 is a group NHCOR¹⁵-or-NHSO₂R¹⁵ where R^{15} is optionally substituted alkyl₇ or optionally substituted aryl-or-optionally substituted heteroaryl;
- R⁵, R⁶ and R⁷ are independently selected from hydrogen, a functional group or an optionally substituted hydrocarbyl group;
- and further provided that when R⁴ is a group NHCOR¹⁵, R¹⁵ is substituted alkyl, optionally substituted aryl or optionally substituted heteroaryl.

- 2. (Previously Presented) A compound according to claim 1 wherein a group R¹⁵ as it appears in the definition of R⁴, is substituted by at least one functional group, or an aryl or heterocyclyl group, either of which may themselves be substituted by one or more functional groups or further aryl or heterocyclyl groups.
- 3. (Previously Presented) A compound according to claim 1 wherein R¹⁵ is a substituted alkyl group or an optionally substituted heterocyclyl or optionally substituted phenyl group.
- 4. (Previously Presented) A compound according to claim 3 wherein R^{15} is alkyl substituted by a group of formula $NR^{19}R^{20}$ where R^{19} and R^{20} are independently selected from hydrogen or optionally substituted hydrocarbyl, or R^{19} and R^{20} together form an optionally substituted ring which optionally contains further heteroatoms such as $S(O)_m$, oxygen and nitrogen, n is an integer of 1 or 2, and m is 1 or 2.
- 5. (Previously Presented) A compound according to claim 1, where R² is carboxy.
- 6. (Previously Presented) A compound according to claim 1 wherein R¹ is 3,4-dichlorophenyl, 3-fluoro-4-chlorophenyl, 3-chloro-4-fluorophenyl or 2,3-dichloropyrid-5-yl.
- 7. (Previously Presented) A compound according to claim 1, where X is CH₂.
- 8. (Currently Amended) A process for preparing a compound according to claim 1, which process comprises reacting a compound of formula (VII)

$$R^{5}$$
 R^{6}
 R^{7}
 R^{1}
 (VII)

where X, R¹, R³, R⁵, R⁶ and R⁷ are as defined in claim 1, and R² is a group R² as defined in relation to formula (I) or a protected form thereof, with a compound of formula (VIII)

 $Z-R^{22}$

(VIII)

where Z is a leaving group and R²² is a group COR¹⁵² or SO₂R¹⁵³ where R¹⁵³ is group R¹⁵ as defined in relation to formula (I) or a precursor thereof; and thereafter if desired or necessary:

- (i) converting a precursor group R^{15} to a group R^{15} and/or converting a group R^{15} to a different such group R^{15} ; and
- (ii) deprotecting a group R2' to a group R2.
- 9. (Previously Presented) A pharmaceutical composition comprising a compound according to claim 1 in combination with a pharmaceutically acceptable carrier.
- 10. (Previously Presented) A method for antagonizing an MCP-1 (Monocyte Chemoattractant Protein-1) or RANTES (Regulated upon Activation, Normal T-cell Expressed and Secreted) mediated effect in a warm blooded animal in need of such treatment comprising administering to said animal an effective amount of a compound according to claim 1, a pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester thereof.
- 11. (Previously Presented) A method for treating inflammation in a warm blooded animal in need of such treatment comprising administering to said animal an effective amount of a compound according to claim 1, a pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester thereof.